AMENDMENTS TO THE CLAIMS

Please amend claims as follow:

1. (Currently Amended) A method of producing a virus comprising:

adhering adhesive cells to a support consisting essentially of nylon, which has a polypeptide of about 20,000 Mn having a structure where 5 (Arg Gly Asp) sequences (SEQ ID NO: 70) and 5 (Gly Ala Gly Ala Gly Ser)₃ sequences (SEQ ID NO: 74) are alternately chemically bonded, and <u>is</u> free from animal-origin components, or a support consisting essentially of nylon, which has a polypeptide of about 10,000 Mn having a structure where 3 (Arg Gly Asp) sequences (SEQ ID NO: 70) and 3 (Gly Val Pro Gly Val)₂ Gly Gly (Gly Ala Gly Ala Gly Ser)₃ sequences (SEQ ID NO: 71) are alternately chemically bonded, and <u>is</u> free from animal-origin components;

culturing the adhesive cells in a medium free from animal-origin components;

subculturing the cultured adhesive cells using a cell dispersing agent that is free from animal-origin components and is a protease originated from a plant, a protease originated from genetically recombinant bacteria, or a combination thereof; and then

inoculating and proliferating a virus in the cells obtained by culturing the adhesive cells, thereby improving efficiency for producing a virus.

- 2. (Previously Presented) The method according to claim 1, wherein said virus belongs to at least one selected from a group consisting of *Flaviviridae*, *Orthomyxoviridae*, *Adenoviridae*, *Herpesviridae*, *Picornaviridae*, *Paramyxoviridae*, *Togaviridae*, and *Poxviridae*.
- 3. (Previously Presented) The method according to claim 1, wherein said support is a microcarrier.
 - 4-6. (Canceled)

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7. (Previously Presented) The method according to claim 2, wherein said support is a microcarrier.

8. (Canceled)